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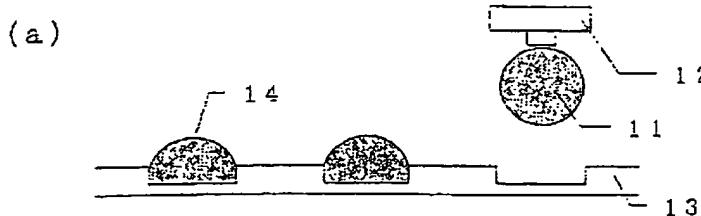
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(54) METHOD FOR FORMING A SILICON FILM AND INK COMPOSITION FOR INK JET

(57) An ink composition 11 containing a silicon precursor is selectively discharged into predetermined regions on a substrate from an ink jet head 12 to form a pattern of the silicon precursor, and is subjected to a heat and/or light treatment to convert the silicon precur-

sor into an amorphous silicon film 15 or a polycrystalline silicon film 16. A silicon film pattern is thereby obtained on a large area at low cost with low energy.

FIG. 1



Description**TECHNICAL FIELD**

5 [0001] The present invention relates to an ink composition containing a silicon precursor for forming by patterning a silicon film by an ink jet process in use of LSIs, thin film transistors, and photosensitive members, and to a method for forming a silicon film by an ink jet process.

BACKGROUND ART

10 [0002] As conventional methods for forming amorphous silicon films and polysilicon films, thermal CVD (chemical vapor deposition) processes, plasma enhanced CVD processes, and photo-assisted CVD processes using monosilane gas and disilane gas have been used. In general, thermal CVD processes are widely used for polysilicon films (refer to J. Vac. Sci. Technology, Vol. 14, 1082 (1977)), and plasma enhanced CVD processes are widely used for amorphous 15 silicon films (refer to Solid State Com., Vol. 17, 1193 (1975)). These are used in production of liquid crystal devices having thin film transistors and solar batteries.

20 [0003] In the formation of silicon films by these CVD processes, however, further improvements are awaited with regard to the following problems. (1) The production yield is low due to system contamination and the formation of foreign materials, which are caused by silicon particles generated during the gas phase reaction. (2) A uniform film thickness is barely obtainable on a substrate having an uneven surface due to gaseous raw materials. (3) The productivity is low due to a low deposition rate of the film. (4) The plasma enhanced CVD process requires a complicated and expensive radiofrequency generator and vacuum equipment.

25 [0004] In regard to materials, use of silicon hydride, which is highly reactive gas, causes difficulty in handling and requires hermetic vacuum equipment. Since these apparatuses are complicated, the apparatuses themselves are expensive. Moreover, the vacuum system and the plasma system consume a large amount of energy, resulting in high production costs.

30 [0005] In recent years, a method is proposed for applying liquid silicon hydride with no vacuum system. Japanese Unexamined Patent Application Publication No. 1-29661 discloses a method for forming a silicon-based thin film in which a gaseous material is liquefied and is adsorbed onto a cooled substrate to react with chemically active, atomic hydrogen. This method, however, has the following problems. (1) Since silicon hydride as the raw material is continuously gasified and are cooled, a complicated apparatus is required and the control of the thickness is difficult.

35 [0006] An example of solid silicon hydride polymers is reported in Great Britain Patent No. GB-2,077,710A. Since this polymer is insoluble in solvents, the film is not formed by coating.

[0007] Moreover, Japanese Unexamined Patent Application Publication No. 9-237927 discloses a method for spray-coating a polysilane solution onto a substrate having a large area and for pyrolyzing the coating to form a silicon film, in order to produce solar batteries. In carbon-containing silicon compounds, a large amount of carbon remains as an impurity in pyrolysis or photolysis by ultraviolet irradiation, hence, it is difficult to prepare an amorphous or polycrystalline silicon film having superior electrical characteristics.

40 [0008] The silicon semiconductor film is generally doped with a Group III element or a Group V element in the Periodic Table and is used as a p-type or n-type semiconductor. Doping using these elements is generally performed by thermal diffusion and ion implantation after the formation of the silicon film. Since the doping is performed *in vacuo*, process control is complicated, and it is particularly difficult to form a uniformly doped silicon film onto a large substrate.

45 [0009] On the other hand, the above Japanese Unexamined Patent Application Publication No. 9-237927 discloses a method for applying a polysilane solution containing an alkyl compound which imparts p-type or n-type conductivity and a method for pyrolyzing a coating film from a polysilane solution in an atmosphere containing a dopant source. In the former method, however, a uniformly doped film may be unobtainable due to differences in solubility between the polysilane and the dopant-containing alkyl compound, and the final film product inevitably contains a large amount of carbon as an impurity due to carbon contained in the material, as described above. In the latter method, it is difficult to control the dose.

50 [0010] In a conventional method for forming a silicon film pattern, a silicon film is formed on the entire substrate by the above method, and a required pattern is formed by a photolithographic process or an etching process. When the silicon film is patterned over a large area to form many devices, many production steps must be employed and expensive apparatuses and various materials are used with a large amount of energy, resulting in high production costs.

DISCLOSURE OF INVENTION

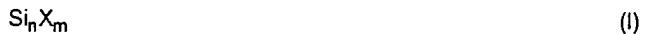
55 [0011] It is an object of the present invention to provide a composition, which is a material for stably forming a desired silicon film pattern with low energy consumption at low cost, and a method for forming a silicon film without a

conventional vacuum process, photolithography, and etching, in the production of a device having a desired silicon film pattern onto a substrate having a large area. It is another object to provide a composition containing a silicon precursor and a method for patterning a silicon film, in the production of a device having a silicon film doped with boron or phosphorus in which a film comprising a modified silicon compound as a silicon precursor is formed using a solution, and
5 the silicon precursor film is converted into semiconductive silicon and doping is simultaneously achieved by heat and/or light treatment in an inert atmosphere.

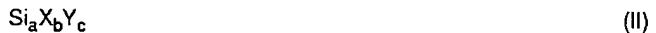
[0012] According to the present invention, there is provided a method for forming a silicon film comprising applying by patterning an ink composition containing a silicon compound onto a substrate by an ink jet process.

[0013] Moreover, the present inventors have intensively researched to achieve the above objects and have been
10 discovered that an ink material composition containing a silicon compound represented by the following general formula (I) or (II) exhibits superior material stability, discharge stability, and coating uniformity in an ink jet process, and that a required silicon film pattern can be formed on a substrate by heat and/or light treatment of a silicon precursor coating film from the solution, with satisfactory conversion into the silicon film without incorporation of a solvent.

[0014] According to the present invention, there is provided a method for a silicon film pattern comprising applying
15 by patterning a composition containing a silicon compound represented by the following general formula (I) or (II) onto a substrate by an ink jet process:



20 wherein n represents an integer 3 or more, m represents an integer of n, 2n-2, 2n, or 2n+2, and X represents a hydrogen atom and/or a halogen atom.



25 wherein X represents a hydrogen atom and/or a halogen atom, Y represents a boron atom or a phosphorus atom, a represents an integer of 3 or more, b represents an integer of a to 2a+c+2, and c represents an integer of 1 to a.

[0015] It is preferable to use a mixture of the silicon compound represented by the general formula (I) and the silicon compound represented by the general formula (II).

[0016] According to the present invention, there is provided a liquid composition containing a silicon compound for
30 ink jet processes comprising a silicon compound represented by the general formula (I) and/or a silicon compound represented by the general formula (II), and a solvent for dissolving the silicon compound, the solvent having a specific viscosity, a specific surface tension, and a specific solid component content.

BRIEF DESCRIPTION OF THE DRAWINGS

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[0017]

Fig. 1 is a drawing showing patterning steps of a silicon thin film by an ink jet process in accordance with Example 1.

40 Fig. 2 is a drawing showing patterning steps of a silicon thin film by an ink jet process in accordance with Example 2.

Fig. 3 is a drawing showing patterning steps of a p-type silicon thin film by an ink jet process in accordance with Example 3.

45 Fig. 4 is a drawing showing patterning steps of an n-type silicon thin film by an ink jet process in accordance with Example 4.

Fig. 5 is a drawing showing patterning steps of a silicon thin film by an ink jet process in accordance with Example 5.

50 Fig. 6 is a drawing showing patterning steps of a p-type silicon thin film by an ink jet process in accordance with Example 6.

Fig. 7 is a drawing showing patterning steps of a silicon thin film by an ink jet process in accordance with Example 7.

BEST MODE FOR CARRYING OUT THE INVENTION

55 [0018] The embodiments of the present invention will be described in detail.

[0019] The silicon compounds used in the method for forming the silicon film in the present invention are preferably compounds represented by the general formula (I) Si_nX_m wherein n represents an integer 3 or more, m represents an integer of n, 2n-2, 2n, or 2n+2, and X represents a hydrogen atom and/or a halogen atom, and compounds represented

by the general formula (II) $\text{Si}_a\text{X}_b\text{Y}_c$ wherein X represents a hydrogen atom and/or a halogen atom, Y represents a boron atom or a phosphorus atom, a represents an integer of 3 or more, b represents an integer of a to $2a+c+2$, and c represents an integer of 1 to a.

[0020] In these general formulae (I) and (II), n represents an integer of 3 or more, and a+c represents an integer of 4 or more. In consideration of thermodynamic stability, solubility, and feasibility of purification, cyclic silicon compounds with n of approximately 5 to 20 and particularly 5 or 6 are preferred in the general formula (I), and silicon compounds with a+c of approximately 5 to 20 and particularly 5 or 6 are preferred in the general formula (II). At a value of less than 5, the silicon compound is unstable due to distortion of the ring and handling is difficult. When n and a+c are larger than 20, the solubility decreases due to cohesive force of the silicon compound and selectable solvents are limited.

[0021] In the general formulae (I) and (II) for the silicon compounds used in the present invention, X represents a hydrogen atom and/or a halogen atom. Since these silicon compounds are precursors for the silicon films, these must be finally converted into amorphous or polycrystalline silicon by heat and/or light treatment. The silicon-hydrogen bonds and the silicon-halogen bonds are cleaved by the above treatment to form silicon-silicon bonds, and are finally converted into silicon. Examples of halogen atoms are a fluorine atom, a chlorine atom, a bromine atom, and an iodine atom. Chlorine and bromine are preferable with regard to the bond cleavage. X may be a hydrogen atom alone or a halogen atom alone, or may be a partially halogenated compound in which the sum of hydrogen atoms and halogen atoms satisfies the general formula (I) or (II).

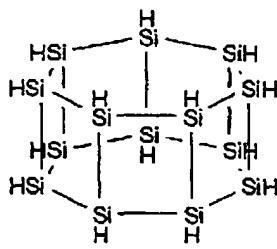
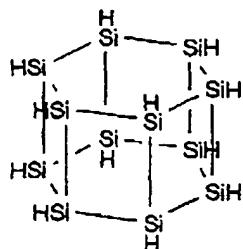
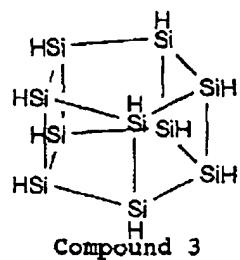
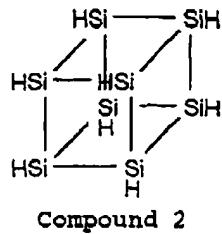
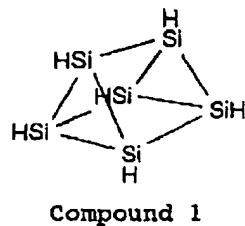
[0022] Examples of the silicon compounds represented by the general formula Si_nX_m used in the present invention will be described. Examples of the compounds of $m = 2n+2$ include silane hydrides, such as trisilane, tetrasilane, pentasilane, hexasilane, and heptasilane, and substituted compounds thereof in which hydrogen atoms are partially or completely replaced with halogen atoms. Examples of $m= 2n$ include monocyclic silicon hydride compounds, such as cyclotrisilane, cyclotetrasilane, cyclopentasilane, silylcyclopentasilane, cyclohexasilane, silylcyclohexasilane, and cycloheptasilane; and halogenated cyclic silicon compounds thereof in which hydrogen atoms are partially or completely replaced with halogen atoms, such as hexachlorocyclotrisilane, trichlorocyclotrisilane, coctachlorocyclotetrasilane, tetrachlorocyclotetrasilane, decachlorocyclopentasilane, pentachlorocyclopentasilane, dodecachlorocyclohexasilane, hexachlorocyclohexasilane, tetradeccachlorocycloheptasilane, heptachlorocycloheptasilane, hexabromocyclotrisilane, tribromocyclotrisilane, pentabromocyclotrisilane, tetrabromocyclotrisilane, octabromocyclotetrasilane, tetrabromocyclotetrasilane, decabromocyclopentasilane, pentabromocyclopentasilane, dodecabromocyclohexasilane, hexabromocyclohexasilane, tetradecabromocycloheptasilane, and heptabromocycloheptasilane. Examples of compounds of $m= 2n-2$ include dicyclic silicon hydride compounds, such as 1,1'-biscyclobutasilane, 1,1'-biscyclopentasilane, 1,1'-biscyclohexasilane, 1,1'-biscycloheptasilane, 1,1'-cyclobutasilylcyclopentasilane, 1,1'-cyclobutasilylcyclohexasilane, 1,1'-cyclobutasilylcycloheptasilane, 1,1'-cyclopentasilylcyclohexasilane, 1,1'-cyclopentasilylcycloheptasilane, spiro[2,2]pentasilane, spiro[3,3]heptasilane, spiro[4,4]nonasilane, spiro[4,5]decasilane, spiro[4,6]undecasilane, spiro[5,5]undecasilane, spiro[5,6]dodecasilane, and spiro[6,6]tridecasilane; substituted silicon compounds in which hydrogen atoms are partly or completely replaced with SiH_3 groups or halogen atoms. Moreover, examples of compounds of $m= n$ include polycyclic silicon hydride compounds, such as Compounds 1 to 5 represented by the following formulae, and substituted silicon compounds thereof in which hydrogen atoms are partially or completely replaced with SiH_3 groups or halogen atoms. These compounds may be used as a mixture of two or more types.

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[0023] Examples of the compounds represented by the general formula (II) $\text{Si}_a\text{X}_b\text{Y}_c$ used in the present invention include modified silane compounds as shown in Compounds 6 to 27 represented by the following formulae.

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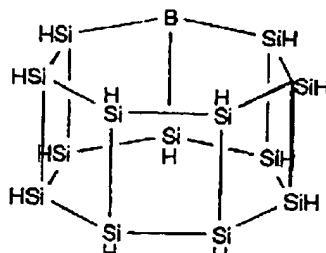
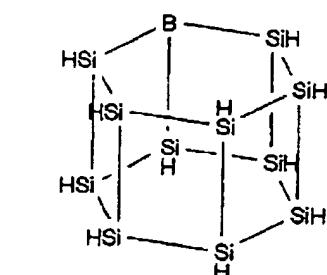
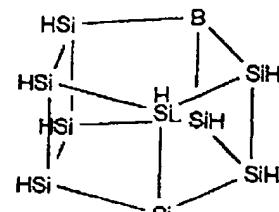
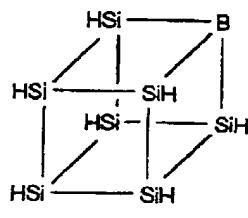
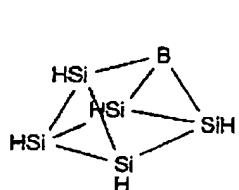
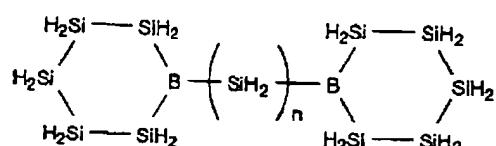
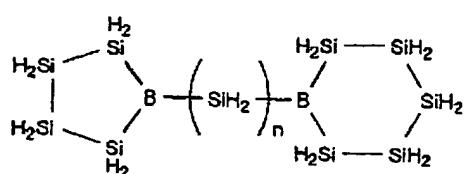
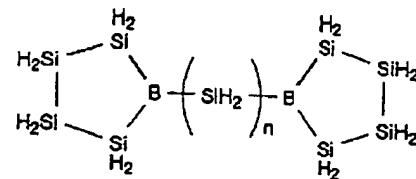
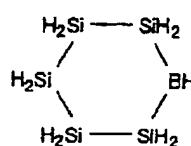
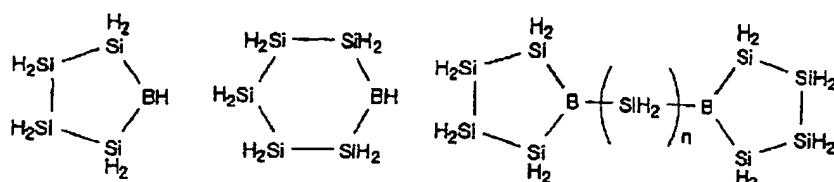
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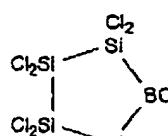
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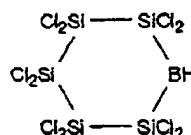


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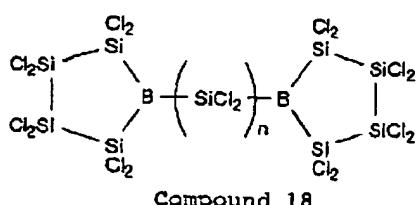
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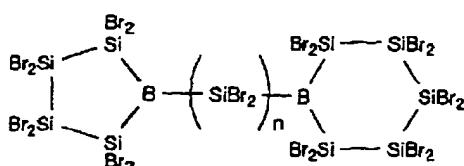
Compound 16



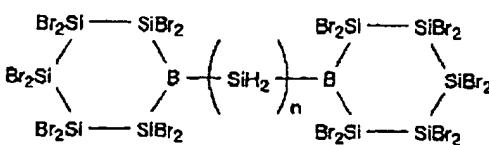
Compound 17



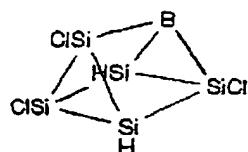
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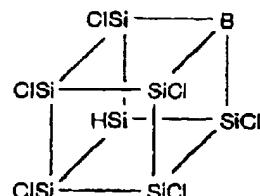
Compound 19



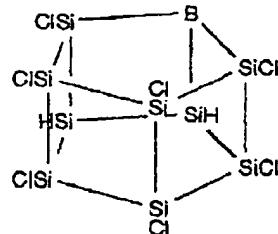
Compound 20



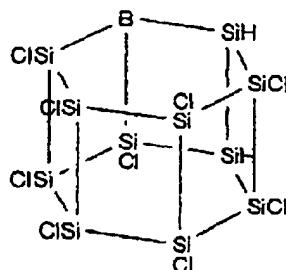
Compound 21



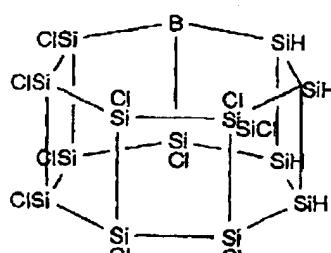
Compound 22



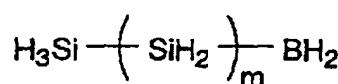
Compound 23



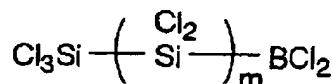
Compound 24



Compound 25



Compound 26



Compound 27

[0024] Herein, n in Compounds 8, 9, 10, 18, 19, and 20 represents an integer of 0 or more, and m in Compounds 26 and 27 represents an integer of 2 or more. In these formulae, boron-modified silane compounds are shown. Phosphorus-modified silane compounds also have similar backbones to those of the boron-modified silane compounds.

[0025] In the present invention, the above modified silane compounds may be used alone as a solution, or may be used as a mixture together with the above unmodified silane compounds. The ratio of the modified silane compound to the unmodified silane compound depends on the boron or phosphorus content as a modifying element, and the modifying element is 1 ppb to 25% to silicon atoms.

5 [0026] In the present invention, a solution in which the above silicon compound dissolved in a solvent is applied to an ink jet process. The vapor pressure of the solvent used in the present invention is preferably in a range of 0.001 to 50 mmHg. A vapor pressure exceeding 50 mmHg readily causes nozzle clogging due to drying of droplets discharged by the ink jet process, and will preclude stable discharge. A vapor pressure of less than 0.001 mmHg causes delayed drying of discharged ink. Since the solvent tends to remain in the silicon compound, a high quality silicon film is barely
10 obtained after the heat and/or light treatment in the subsequent step.

[0027] Solvents used in the present invention are not limited as long as these can dissolve the silicon compounds and are unreactive with the compounds. Examples of the solvents include hydrocarbon solvents, such as n-heptane, n-octane, decane, toluene, xylene, cymene, durene, indene, dipentene, tetrahydronaphthalene, decahydronaphthalene, and cyclohexylbenzene; ether solvents, such as ethylene glycol dimethyl ether, ethylene glycol diethyl ether, ethylene
15 glycol methyl ethyl ether, diethylene glycol dimethyl ether, diethylene glycol diethyl ether, diethylene glycol methyl ethyl ether, 1,2-dimethoxyethane, bis(2-methoxyethyl)ether, and p-dioxane; and polar solvents, such as propylene carbonate, γ -butyrolactone, N-methyl-2-pyrrolidone, dimethylformamide, dimethyl sulfoxide, and cyclohexanone. Among these, hydrocarbon solvents and ether solvents are preferable in view of solubility of silicon compounds and stability thereof. The hydrocarbon solvents are more preferable. These solvents may be used alone or in combinations of two or
20 more. In particular, the hydrocarbon solvents are advantageous, since these exhibit high solubility of the silicon compounds and suppress the remaining silicon compound after the heat treatment and the light treatment, described below.

[0028] In the present invention, the ink composition (solution) applied to the ink jet process is prepared by mixing the above silicon compounds and/or the modified silicon compounds as solutes with the above solvents. The concentration of the solutes are determined depending on the required thickness of the silicon film. Preferably, the concentration
25 is 0.01 to 10 percent by weight. When the concentration is less than 0.01%, a uniform coating film is not obtained and discharge operations must be increased. When the concentration exceeds 10%, the viscosity increases and affects discharge.

[0029] These solvents may contain a surface tension modifier, such as a fluorine type, a silicon type, or a nonionic type, in a trace amount, if necessary, within a range which does not impair required functions. These solvents may contain a surface tension modifier, such as a fluorine type, a silicon type, or a nonionic type, in a trace amount, if necessary, within a range which does not impair required functions. The nonionic surface tension modifier improves wettability to a workpiece for applying the solution and leveling of the coating film, and prevents graininess and an orange peel surface of the coating film.

[0030] Examples of such nonionic surfactants include fluorinated surfactants having fluoroalkyl groups or perfluoroalkyl groups, and polyether alkyl surfactants having oxyalkyl groups. Examples of fluorinated surfactants include C₉F₁₉CONHC₁₂H₂₅, C₈F₁₇SO₂NH-(C₂H₄O)₆H, C₉F₁₇O-(PLURONIC L-35)-C₉F₁₇, C₉F₁₇O-(PLURONIC P-84)-C₉F₁₇, and C₉F₇O-(TETRONIC-704)-(C₉F₁₇)₂, wherein PLURONIC L-35 is a polyoxypropylene-polyoxyethylene block copolymer having an average molecular weight of 1,900 made by Asahi Denka Kogyo K.K., PLURONIC P-84 is a polyoxypropylene-polyoxyethylene block copolymer having an average molecular weight of 4,200 made by Asahi Denka Kogyo K.K., and TETRONIC-704 is an N,N,N',N'-tetrakis(polyoxypropylene-polyoxyethylene) block copolymer having an average molecular weight of 5,000 made by Asahi Denka Kogyo K.K.

[0031] Actual examples of the fluorinated surfactants include Efttop EF301, Efttop EF303, and Efttop EF352 made by Shin Akita Kasei K.K.; Megafack F171 and Megafac F173 made by Dainippon Ink and Chemicals; Asahi Guard AG710 made by Asahi Glass Co., Ltd.; Fluorad FC-170C, Fluorad FC430, and Fluorad FC431 made by Sumitomo 3M Limited; Surflon S-382, Surflon SC101, Surflon SC102, Surflon SC103, Surflon SC104, Surflon SC105, and Surflon SC106 made by Asahi Glass Co., Ltd.; BM-1000 and BM-1100 made by B. M-Chemie; and Schsego-Fluor made by Schwemann.

[0032] Polyether alkyl surfactants are polyoxyethylene alkyl ethers, polyoxyethylene allyl ether, polyoxyethylene alkylphenol ethers, polyoxyethylene fatty acid esters, sorbitan fatty acid esters, polyoxyethylene sorbitan fatty acid ethers, and oxyethylene-oxypropylene block copolymers.

[0033] Actual examples of these polyether alkyl surfactants include Emulgen 105, Emulgen 430, Emulgen 810, Emulgen 920, Rheodol SP-40S, Rheodol TW-L120, Emanol 3199, Emanol 4110, Excel P-40S, Bridge 30, Bridge 52, Bridge 72, Bridge 92, Arlacel 20, Emasol 320, Twin 20, Twin 60, and Merge 45 made by Kao Corporation; and Nonibol 55 made by Sanyo Chemical Industries, Ltd. Examples of other nonionic surfactants include polyoxyethylene fatty acid esters, polyoxyethylene sorbitan fatty acid esters, and polyalkylene oxide block copolymers. Actual Examples of these surfactants are Chemistat 2500 made by Sanyo Chemical Industries, Ltd., SN-EX9228 made by San Nopco Limited, and Nonal 530 made by Toho Chemical Industry Co., Ltd.

[0034] It is preferable that the viscosity of the prepared silicon-compound-containing solution be within a range of

1 to 50 mPa·s. When the viscosity is lower than 1 mPa·s, the peripheral section of the nozzle in the ink jet process is readily contaminated due to overflow of the ink. When the viscosity is higher than 50 mPa·s, the frequency of nozzle clogging increases to preclude smooth discharge of droplets.

[0035] Preferably, the surface tension of the silicon-compound-containing solution is in a range of 20 to 70 dyn/cm.

5 A surface tension of less than 20 dyn/cm causes increased wettability of the ink composition to the nozzle surface, resulting in nonlinear flight. A surface tension exceeding 70 dyn/cm causes an unstable meniscus shape at the nozzle tip, resulting in uncontrollability of the discharged volume and discharging timing of the ink composition.

[0036] The droplet discharge apparatus in the ink jet process used for supplying the solution in the present invention may be of any mechanism which can discharge appropriate droplets in constant volumes. The apparatus may be 10 of an ink jet process using a piezoelectric element which can form droplets, each weights approximately several tens nanograms, or of a bubble jet process which forms bubbles by thermal energy from a heater. Moreover, a general coating process, such as spin coating, dip coating, spray coating, roll coating, or curtain coating, may be combined, if necessary. When the ink composition containing the silicon precursor is applied by the ink jet process, the atmosphere is generally an inert gas, such as argon, helium, or nitrogen, and the temperature is appropriately selected in a range of 15 room temperature to approximately 100°C depending on the ink characteristics.

[0037] In the present invention, the substrate used in discharge of the silicon-compound-containing solution by the ink jet process is not limited. Examples of the substrates include general quartz, borosilicate glass, soda glass, transparent electrodes such as ITO, metallic substrates, e.g., gold, silver, copper, nickel, titanium, aluminum, and tungsten, and glass or plastic substrates coated with these metals.

20 [0038] The coating film of the silicon-compound-containing solution formed by the ink jet process is subjected to a heat and/or light treatment. The treatment is performed in an inert gas atmosphere, such as nitrogen, argon, and helium. A small amount of reducing gas such as hydrogen may be added, if necessary. The heat treatment is performed for the removal of the solvent and the conversion into a metallic silicon film, after the coating of the ink (solution). The drying treatment may be appropriately determined by the boiling point (vapor pressure) and the pressure of the solvent 25 and thermal behavior of the silicon compound, without particular limitation. The conversion into the metallic silicon film is generally performed in an argon atmosphere at approximately 100 to 800°C, preferably approximately 200 to 600°C, and more preferably 300°C to 500°C. In general, an amorphous silicon film is obtained at a final temperature of 550°C or less, whereas a polycrystalline silicon film is obtained at a higher temperature. At a final temperature of less than 300°C, pyrolysis of the silicon compound does not sufficiently proceed, and a silicon film having a sufficient thickness 30 cannot be formed in some cases. The polycrystalline silicon film may also be prepared by conversion of the above amorphous silicon film into the polycrystalline silicon by laser irradiation. The atmosphere in the laser irradiation is preferably an inert gas, such as helium and argon, and may contain a reducing gas such as hydrogen.

[0039] The coating film of the silicon compound formed by the ink jet process in the present invention may be subjected to a light treatment in an inert gas atmosphere before and/or after removing the solvent in the coating film. The 35 silicon compound soluble in solvents is converted into a tough coating film, which is insoluble in the solvents, by the ring cleavage reaction by means of this light treatment. Moreover, the heat treatment, which is performed after the light treatment or simultaneously performed with the light treatment, converts the silicon film into one having superior optical and electrical characteristics.

[0040] Examples of light sources used in the present invention include low-pressure or high-pressure mercury 40 lamps, deuterium lamps, discharge light of rare gases, such as argon, krypton, and xenon, YAG lasers, argon lasers, carbon dioxide lasers, excimer lasers of XeF, XeCl, XeBr, KrF, KrCl, ArF, and ArCl. These light sources generally have power in a range of 10 to 5,000 W, and a power of 100 to 1,000 W is adequate in the present invention. The wavelengths of these light sources are not limited as long as the light is absorbed in the coating film of the photo-ring-cleavage silicon compound to some extents, and are generally in a range of 170 nm to 600 nm. The use of laser light is particularly preferable in view of conversion efficiency into the polycrystalline silicon film. The temperature for the light treatment is generally in a range of room temperature to 500°C, and can be appropriately determined depending on the semiconductive characteristics of the resulting silicon film.

[0041] The present invention will now be described in more detail with reference to the Examples, but should not be limited to these Examples.

50 [0042] In these Examples, the viscosity and the surface tension of each ink composition indicate values at 20°C.

(EXAMPLE 1)

[0043] A silicon-compound-containing solution (ink) was prepared by dissolving 0.05 g of cyclopentasilane into a 55 mixed solvent of 7 g of decahydronaphthalene and 3 g of tetrahydronaphthalene. The viscosity of the solution was 3.2 mPa·s and the surface tension was 33 dyn/cm. Fig. 1 shows the patterning of silicon thin films by an ink jet process in accordance with this example. The silicon-compound-containing ink 11 was pattern-coated onto a quartz substrate 13 using an ink head 12 composed of a piezoelectric element in an argon atmosphere. The substrate was dried at

200°C in an argon atmosphere and was pyrolyzed at 500°C to form a pattern of metallic silicon films 15 (film regions, each having a diameter of 50 µm), each having a thickness of 35 nm, shown in Figs. 1(b) and 1(c). Fig. 1(b) is a cross-sectional view and Fig. 1(c) is a top view corresponding to Fig. 1(b). A surface analysis of the silicon film by ESCA detected only silicon atoms. The crystal state of the silicon film by a Raman spectrum was 100% amorphous. The amorphous silicon films were irradiated with an excimer laser having a wavelength of 308 nm and an energy density of 300 mJ/cm² in an argon atmosphere to form a pattern of polycrystalline silicon films 16 (film regions, each having a diameter of 50 µm) shown in Fig. 1(d). The Raman spectrum showed that the crystallinity was 80%.

(EXAMPLE 2)

[0044] A silicon-compound-containing solution (ink) was prepared by dissolving 0.2 g of hexachlorocyclohexasilane into a mixed solvent of 5 g of cyclohexylbenzene and 5 g of cymene. The viscosity of the solution was 4.8 mPa·s and the surface tension was 32 dyn/cm. Fig. 2 shows the patterning of silicon thin films by an ink jet process in accordance with this example. The silicon-compound-containing ink 21 was pattern-coated onto a quartz substrate 23 using an ink jet head 22 composed of a piezoelectric element in an argon atmosphere. The substrate was dried at 250°C in an argon atmosphere containing 5% hydrogen and was heat-treated at 450°C while being irradiated with ultraviolet light from a 500-W high-pressure mercury lamp for 30 minutes to form a pattern of metallic silicon films 25 (film regions, each being 50 × 50 µm), each having a thickness of 80 nm, shown in Figs. 2(b) and 2(c). A surface analysis of the silicon film by ESCA detected only silicon atoms. The crystal state of the silicon film by a Raman spectrum was 100% amorphous.

The amorphous silicon film was heat-treated at 800°C in an argon/hydrogen atmosphere to form a pattern of a polycrystalline silicon film 26 (film region of 50 × 50 µm) having a crystallinity of 50%, shown in Fig. 2(d).

(EXAMPLE 3)

[0045] A silicon-compound-containing solution (ink) was prepared by dissolving 0.1 g of 1-borahexaprismane (Compound 11) into a mixed solvent of 7 g of tetrahydronaphthalene and 3 g of cyclohexylbenzene. The viscosity of the solution was 4.2 mPa·s and the surface tension was 36 dyn/cm. Fig. 3 shows the patterning of silicon thin films by an ink jet process in accordance with this example. The silicon-compound-containing ink 31 was pattern-coated onto a quartz substrate 33 using an ink jet head 32 composed of a piezoelectric element in an argon atmosphere. The substrate was dried at 250°C in an argon atmosphere and was pyrolyzed at 500°C to form a pattern of metallic silicon films 35 (film regions, each having a diameter of 50 µm), each having a thickness of 70 nm, shown in Figs. 3(b) and 3(c). A surface analysis of the silicon film by ESCA detected only silicon atoms and boron atoms and the ratio thereof was 5:1. The crystal state of the silicon film by a Raman spectrum was 100% amorphous. The amorphous silicon films were irradiated with an excimer laser having a wavelength of 308 nm and an energy density of 350 mJ/cm² in an argon atmosphere to form a pattern of polycrystalline silicon films 36 (film regions, each having a diameter of 50 µm) shown in Fig. 3(d). The Raman spectrum showed that the crystallinity was 75%.

(EXAMPLE 4)

[0046] A silicon-compound-containing solution (ink) was prepared by dissolving 0.05 g of 1-phosphocyclopentasilane into a mixed solvent of 8 g of decahydronaphthalene and 2 g of cyclohexylbenzene. The viscosity of the solution was 4.0 mPa·s and the surface tension was 36 dyn/cm.

[0047] Fig. 4 shows the patterning of silicon thin films by an ink jet process in accordance with this example. The silicon-compound-containing ink 41 was pattern-coated onto a quartz substrate 43 using an ink jet head 42 composed of a piezoelectric element in an argon atmosphere. The substrate was dried at 150°C in an argon atmosphere containing 5% hydrogen while being irradiated with ultraviolet light from a 500-W high-pressure mercury lamp for 10 minutes to form a tough coating film of polysilazane which was insoluble in solvents by a ring cleavage reaction. The coating film was pyrolyzed at 500°C in an argon atmosphere to form a pattern of metallic silicon films 45 (film regions, each being 50 × 50 µm), each having a thickness of 25 nm, shown in Figs. 4(b) and 4(c). A surface analysis of the silicon film by ESCA detected only silicon atoms and phosphorus atoms and the ratio thereof was 5:1. The crystal state of the silicon film by a Raman spectrum was 100% amorphous. The amorphous silicon films were irradiated with an excimer laser having a wavelength of 308 nm and an energy density of 300 mJ/cm² in an argon atmosphere to form a pattern of polycrystalline silicon films 46 (film regions, each being 50 × 50 µm) shown in Fig. 4(d). The Raman spectrum showed that the crystallinity was 75%.

(EXAMPLE 5)

[0048] A silicon-compound-containing solution (ink) was prepared by dissolving 0.5 g of silylcyclopentasilane into

a mixed solvent of 5 g of decahydronaphthalene and 5 g of tetrahydronaphthalene. The viscosity of the solution was 5.2 mPa·s and the surface tension was 34 dyn/cm. Fig. 5 shows the patterning of silicon thin films by an ink jet process in accordance with this example. The silicon-compound-containing ink 51 was pattern-coated onto a polyimide film substrate 53 using an ink jet head 52 composed of a piezoelectric element in an argon atmosphere. The substrate was dried at 200°C in an argon atmosphere and was irradiated using a 500-W high-pressure mercury lamp at 350°C for photolysis and pyrolysis to form a pattern of metallic silicon films 55 (film regions, each having a diameter of 70 µm), each having a thickness of 80 nm, shown in Fig. 5(b). A surface analysis of the silicon film by ESCA detected only silicon atoms. The crystal state of the silicon film by a Raman spectrum was 100% amorphous.

10 (EXAMPLE 6)

[0049] A (p-type) silicon-compound-containing solution (ink) was prepared by dissolving 0.01 g of 1-boracyclopentasilane and 0.1 g of cyclopentasilane into a mixed solvent of 5 g of tetrahydronaphthalene and 5 g of cyclohexylbenzene. The viscosity of the solution was 3.8 mPa·s and the surface tension was 35 dyn/cm. Next, a (n-type) silicon-compound-containing solution (ink) was prepared by dissolving 0.01 g of 1-phosphocyclopentasilane and 0.1 g of cyclopentasilane into a mixed solvent of 5 g of tetrahydronaphthalene and 5 g of cyclohexylbenzene. The viscosity of the solution was 3.7 mPa·s and the surface tension was 36 dyn/cm. Fig. 6 shows the patterning of n-type and p-type silicon thin films by an ink jet process in accordance with this example. The (n-type) silicon-compound-containing ink 61 and the (p-type) silicon-compound-containing ink 64 were pattern-coated onto a quartz substrate 63 using an ink jet head 62 composed of a piezoelectric element in an argon atmosphere. The substrate was dried at 250°C in an argon atmosphere containing 5% hydrogen and was pyrolyzed at 500°C to form a pattern including n-type silicon films 65n (film regions, each being 50 × 50 µm) and p-type silicon films 65p (film regions, each being 50 × 50 µm) shown in Figs. 6(b) and 6(c). According to a surface analysis of the silicon films by ESCA, the ratio of silicon atoms and phosphorus atoms was 50:1 in the n type and the ratio of silicon atoms and phosphorus atoms was 50:1 in the p type. The crystal state of the silicon films by a Raman spectrum was 100% amorphous. The amorphous silicon films were irradiated with an excimer laser having a wavelength of 308 nm and an energy density of 320 mJ/cm² in an argon atmosphere to form a pattern of n-type and p-type polycrystalline silicon films 66n and 66p, respectively, (film regions, each being 50 × 50 µm) shown in Fig. 6(d). The Raman spectrum showed that the crystallinity was 75%.

30 (EXAMPLE 7)

[0050] A silicon-compound-containing solution (ink) was prepared by dissolving 0.05 g of octasilacubane (Compound 2) into a mixed solvent of 5 g of decahydronaphthalene and 5 g of tetrahydronaphthalene. The viscosity of the solution was 5.5 mPa·s and the surface tension was 34 dyn/cm. Fig. 7 shows the patterning of silicon thin films by an ink jet process in accordance with this example. The silicon-compound-containing ink 71 was pattern-coated onto a quartz substrate 73 using an ink jet head 72 composed of a piezoelectric element in an argon atmosphere containing 3% hydrogen. The substrate was irradiated with using a high-pressure mercury lamp of 500 W at room temperature for 5 minutes in an argon atmosphere containing 3% hydrogen. As a result, polyhydrosilane films, which were insoluble in solvents, were formed. The films were pyrolyzed in the argon atmosphere containing 3% hydrogen at 500°C to form a pattern of metallic silicon films 75 (film regions, each having a diameter of 50 µm), each having a thickness of 35 nm, shown in Figs. 7(b) and 7(c). Fig. 7(b) is a cross-sectional view and Fig. 7(c) is a top view corresponding to Fig. 7(b). A surface analysis of the silicon film by ESCA detected only silicon atoms. The crystal state of the silicon film by a Raman spectrum was 100% amorphous. The amorphous silicon films were irradiated with an excimer laser having a wavelength of 308 nm and an energy density of 300 mJ/cm² in an argon atmosphere to form a pattern of polycrystalline silicon films 76 (film regions, each having a diameter of 50 µm) shown in Fig. 7(d). The Raman spectrum showed that the crystallinity was 80%.

(Example 8)

50 [0051] The silicon-compound-containing solution of Example 7 was prepared and was pattern-coated onto a quartz substrate using an ink jet head composed of a piezoelectric element in an argon atmosphere containing 3% hydrogen. The substrate was dried at 150°C in an argon atmosphere containing 3% hydrogen, and simultaneously was irradiated using a 500 W high-pressure mercury lamp for 5 minutes. As a result, polyhydrosilane films, which were insoluble in solvents, were formed. The films were pyrolyzed as in Example 7 to form a pattern of amorphous metallic silicon films, as in Example 7. The amorphous silicon films were irradiated with an excimer laser having the wavelength and the same energy density as those in Example 7 in air to form a pattern of polycrystalline silicon films 76 (film regions, each having a diameter of 50 µm) shown in Fig. 7(d). The Raman spectrum showed that the crystallinity was 60%.

[0052] According to the present invention as described above, a composition containing a silicon precursor is pat-

tern-coated by an ink jet process, and a desired silicon film (pattern) as an electronic material can be formed by a novel liquid phase process, which is different from conventional methods for forming silicon film, by means of thermal and/or optical energy.

5 INDUSTRIAL APPLICABILITY

[0053] Since the present invention prevents the generation of powder during the formation of the silicon film and does not use a large vacuum process, unlike conventional CVD processes, the present invention does not require expensive equipment and can form a film on a substrate having a large area using minimum amounts of materials.

10 Since the silicon film pattern can be formed without a photolithographic etching process, semiconductor devices having silicon films, such as LSIs, thin-film transistors, photoelectric transducers, and photosensitive members, at reduced costs and by a low energy process.

Claims

15 1. A method for forming a silicon film comprising applying by patterning an ink composition containing a silicon compound onto a substrate by an ink jet process.

20 2. A method for forming a silicon film according to claim 1, wherein the ink composition is applied in an inert atmosphere.

25 3. A method for forming a silicon film according to either claim 1 or 2, further comprising, after applying the ink composition containing the silicon compound onto the substrate by the ink jet process, a drying step of removing a solvent of the composition and a step of pyrolyzing and/or photolyzing in the coating film.

30 4. A method for forming a silicon film according to claim 3, further comprising a step for irradiating the silicon film formed by the heat treatment and/or the light treatment with laser to convert the amorphous silicon film into a polycrystalline silicon film.

35 5. A method for forming a silicon film according to claim 1, wherein the silicon compound is a silicon compound represented by the following general formula (I):



35 wherein n represents an integer of 3 or more, m represents an integer of n, 2n-2, 2n, or 2n+2, and X represents a hydrogen atom and/or a halogen atom.

40 6. A method for forming a silicon film according to claim 1, wherein the silicon compound is a silicon compound represented by the following general formula (II):

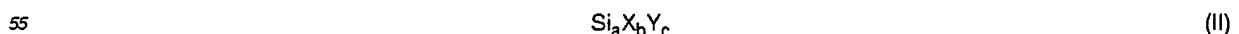


45 wherein X represents a hydrogen atom and/or a halogen atom, Y represents a boron atom or a phosphorus atom, a represents an integer of 3 or more, b represents an integer of a to 2a+c+2, and c represents an integer of 1 to a.

45 7. A method for forming a silicon film according to claim 1, wherein the silicon compound is a composition containing a silicon compound represented by the following general formula (I) and a silicon compound represented by the following general formula (II):



50 wherein n represents an integer 3 or more, m represents an integer of n, 2n-2, 2n, or 2n+2, and X represents a hydrogen atom and/or a halogen atom;



55 wherein X represents a hydrogen atom and/or a halogen atom, Y represents a boron atom or a phosphorus atom, a represents an integer of 3 or more, b represents an integer of a to 2a+c+2, and c represents an integer of 1 to a.

8. A method for forming a silicon film according to either claim 5 or 7, wherein n is in a range of 5 to 20 in the general formula (I).

5 9. A method for forming a silicon film according to either claim 6 or 7, wherein a+c is in a range of 5 to 20 in the general formula (II).

10. A method for forming a silicon film according to any one of claims 1 to 9, wherein, in the composition, the silicon compound is dissolved in at least one solvent having a vapor pressure at room temperature of 0.001 to 50 mmHg.

11. A method for forming a silicon film according to claim 10, wherein the solvent is a hydrocarbon solvent.

12. A method for forming a silicon film according to any one of claims 1 to 11, wherein the concentration of the silicon compound in the composition is in a range of 0.01 to 10 percent by weight.

15 13. A method for forming a silicon film according to any one of claims 1 to 12, wherein the concentration has a viscosity of 1 to 50 mPa · s and a surface tension of 20 to 70 dyn/cm.

14. An ink-jet ink composition for forming a silicon film, comprising a silicon compound represented by the general formula (I):



wherein n represents an integer 3 or more, m represents an integer of n, 2n-2, 2n, or 2n+2, and X represents a hydrogen atom and/or a halogen atom.

25 15. An ink-jet ink composition for forming a silicon film, comprising a silicon compound represented by the general formula (II):



wherein X represents a hydrogen atom and/or a halogen atom, Y represents a boron atom or a phosphorus atom, a represents an integer of 3 or more, b represents an integer of a to 2a+c+2, and c represents an integer of 1 to a.

35 16. An ink-jet ink composition for forming a silicon film, comprising a silicon compound represented by the following general formula (I) and a silicon compound represented by the following general formula (II):



wherein n represents an integer 3 or more, m represents an integer of n, 2n-2, 2n, or 2n+2, and X represents a hydrogen atom and/or a halogen atom;



wherein X represents a hydrogen atom and/or a halogen atom, Y represents a boron atom or a phosphorus atom, a represents an integer of 3 or more, b represents an integer of a to 2a+c+2, and c represents an integer of 1 to a.

50 17. An ink composition according to either claim 14 or 16, wherein n is in a range of 5 to 20 in the general formula (I).

18. An ink composition according to either claim 15 or 16, wherein a+c is in a range of 5 to 20 in the general formula (II).

55 19. An ink composition according to any one of claims 14 to 18, wherein the silicon compound is dissolved in at least one solvent having a vapor pressure at room temperature of 0.001 to 50 mmHg.

20. An ink composition according to claim 19, wherein the solvent is a hydrocarbon solvent.

55 21. An ink composition according to any one of claims 14 to 20, wherein the concentration of the silicon compound in the composition is in a range of 0.01 to 10 percent by weight.

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22. An ink composition according to any one of claims 14 to 21, wherein the concentration has a viscosity of 1 to 50 mPa·s and a surface tension of 20 to 70 dyn/cm.

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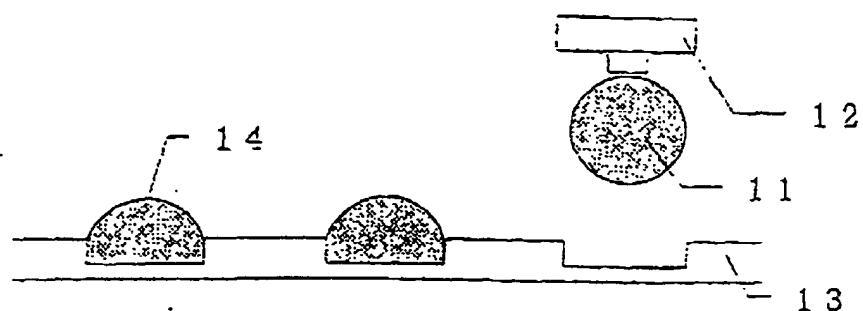
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FIG. 1

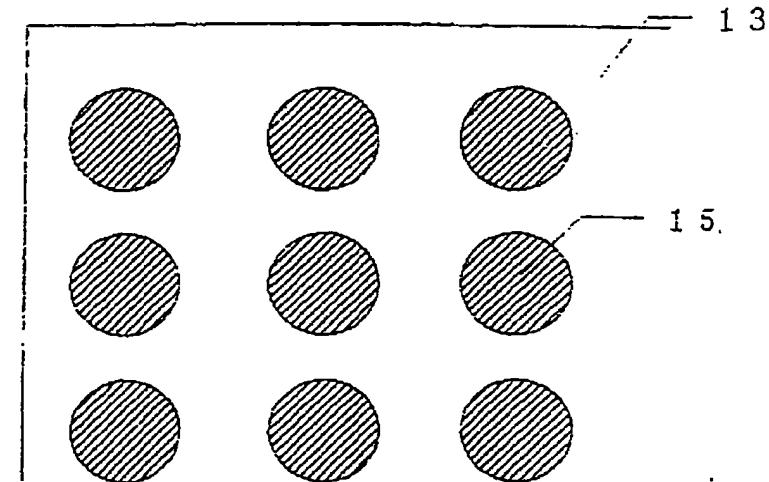
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(d)

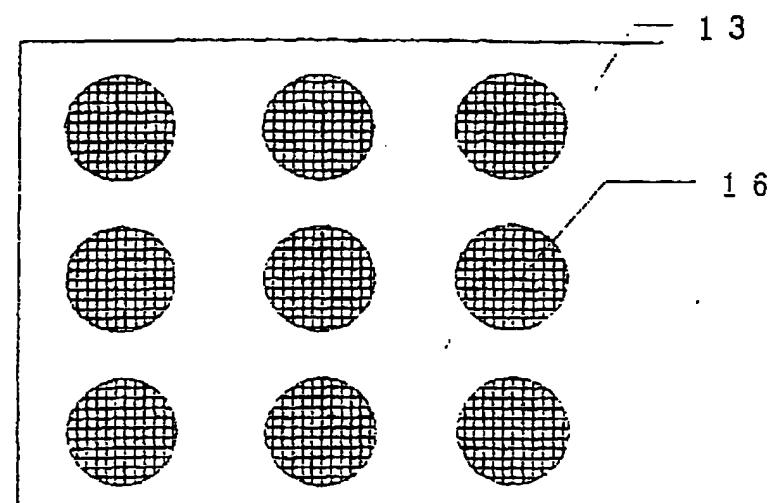
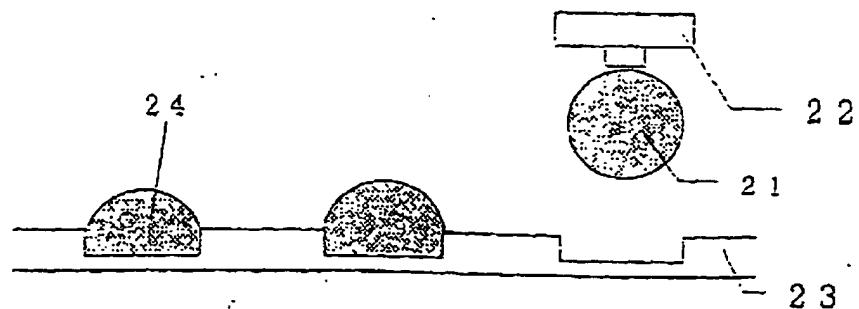
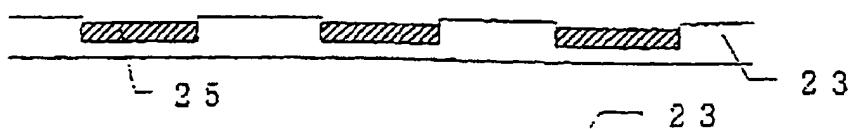


FIG. 2

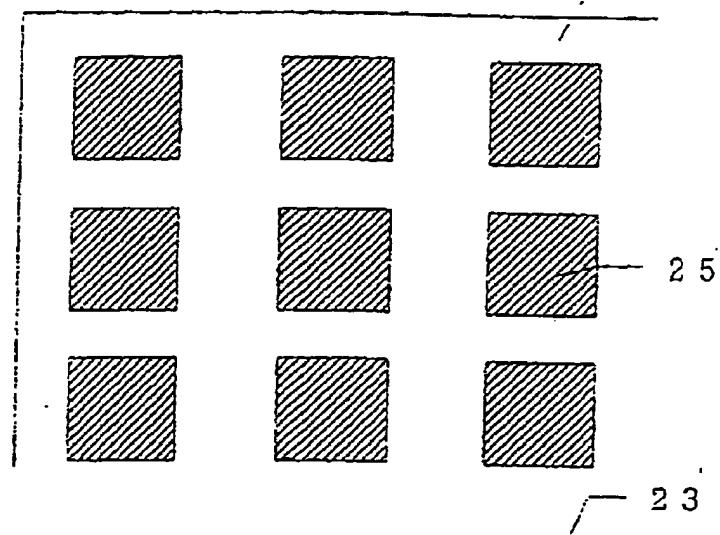
(a)



(b)



(c)



(d)

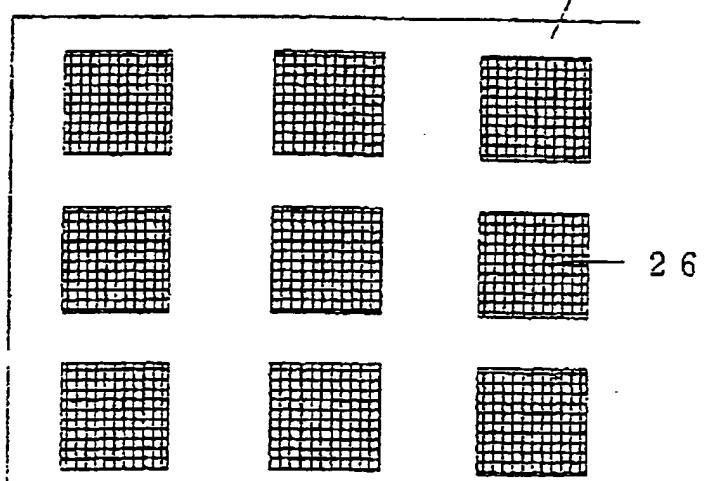
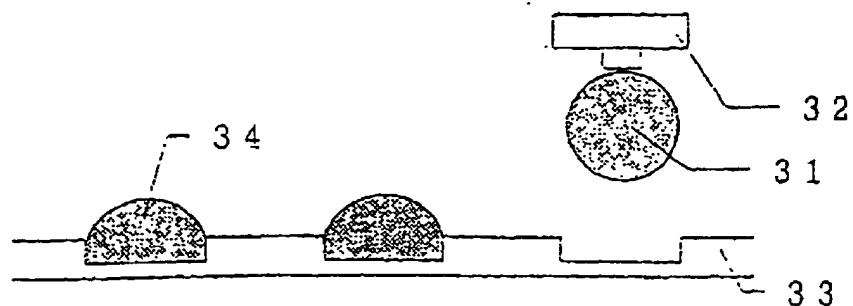
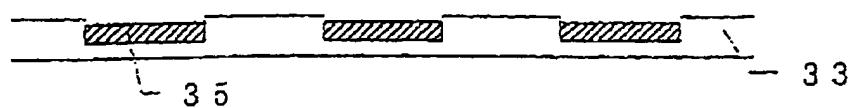


FIG. 3

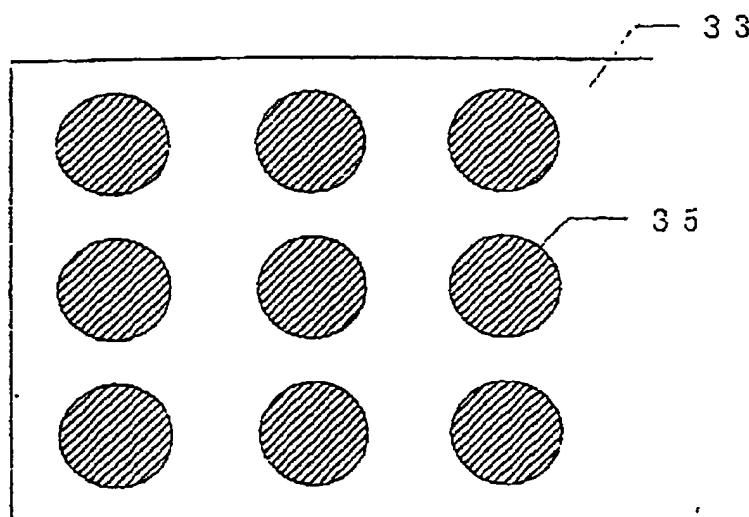
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(b)



(c)



(d)

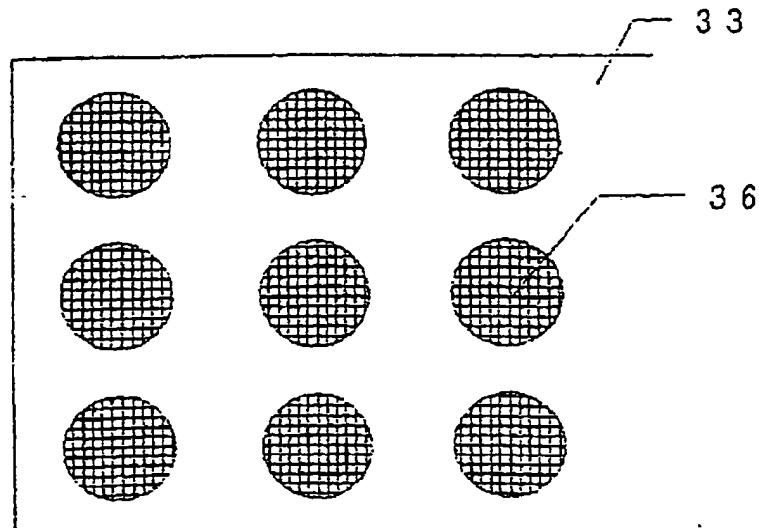
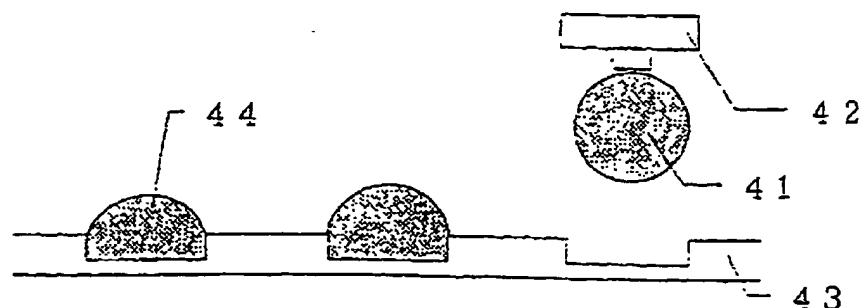
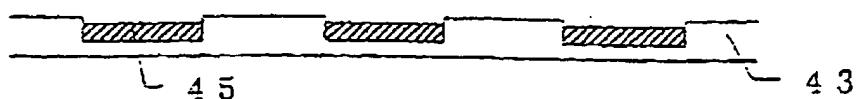


FIG. 4

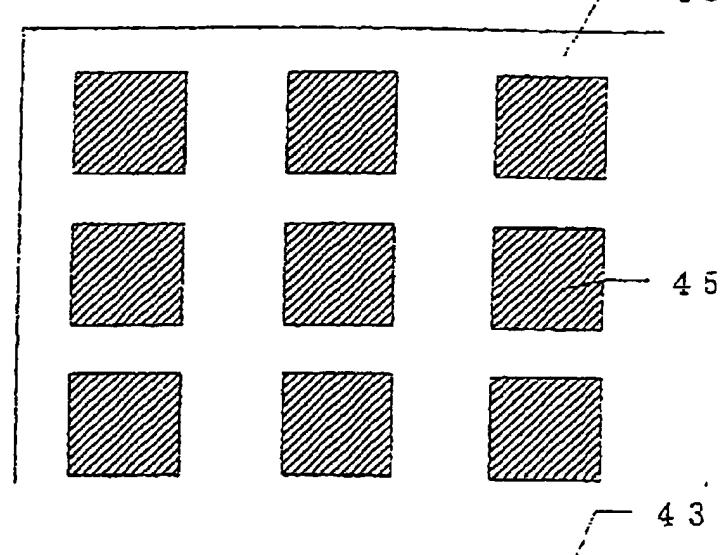
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(d)

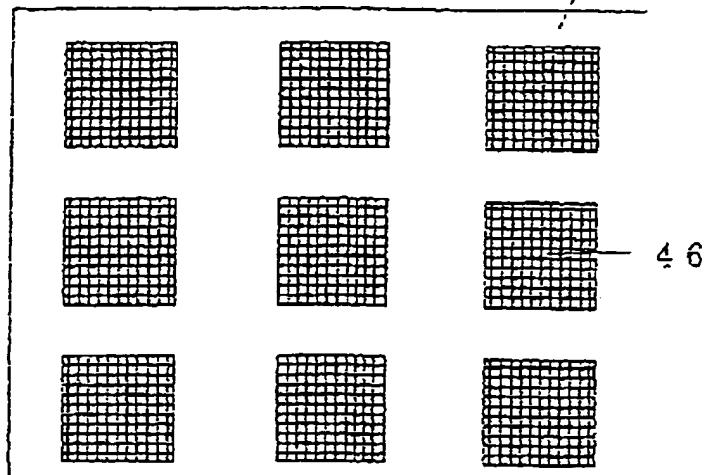
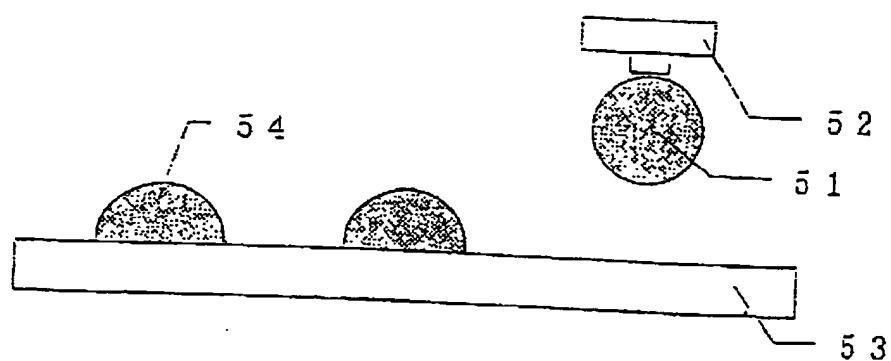


FIG. 5

(a)



(b)

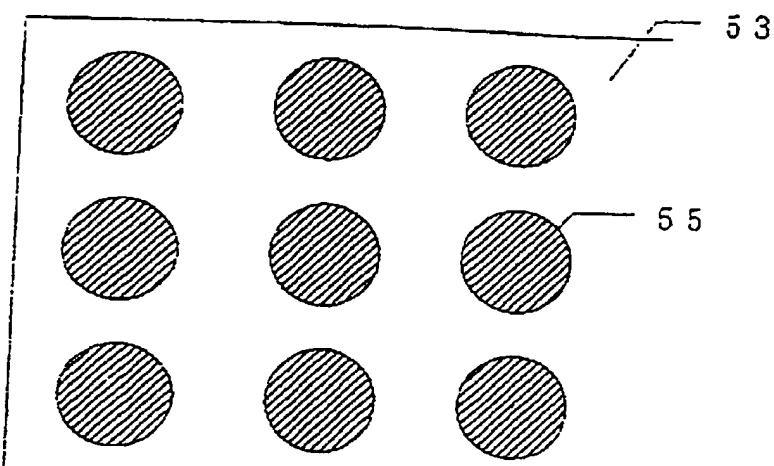
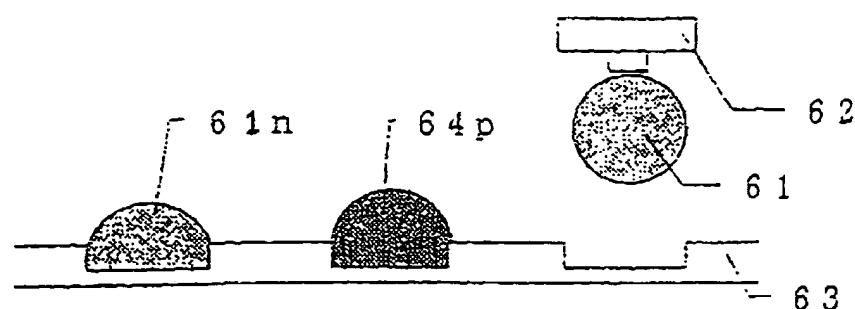
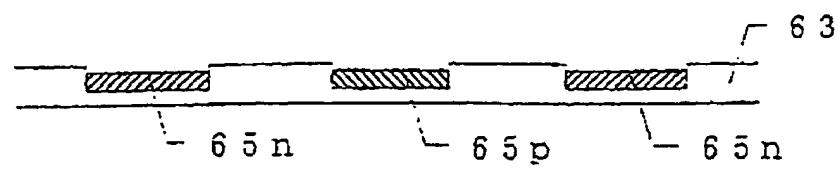


FIG. 6

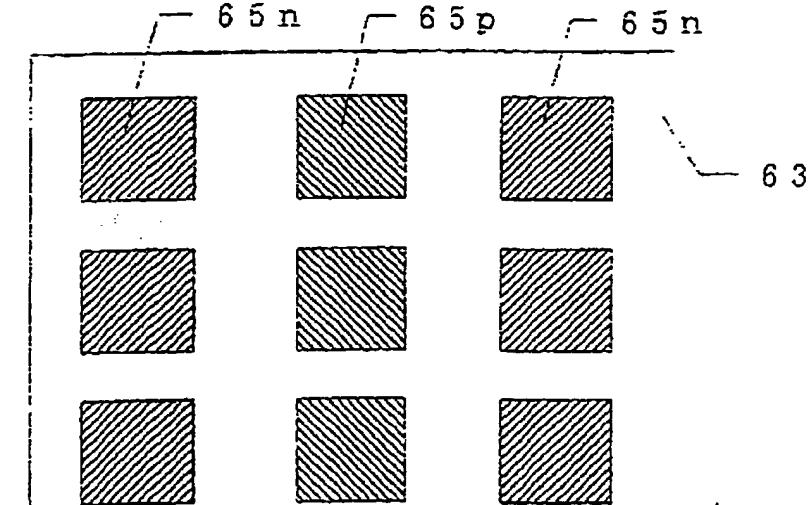
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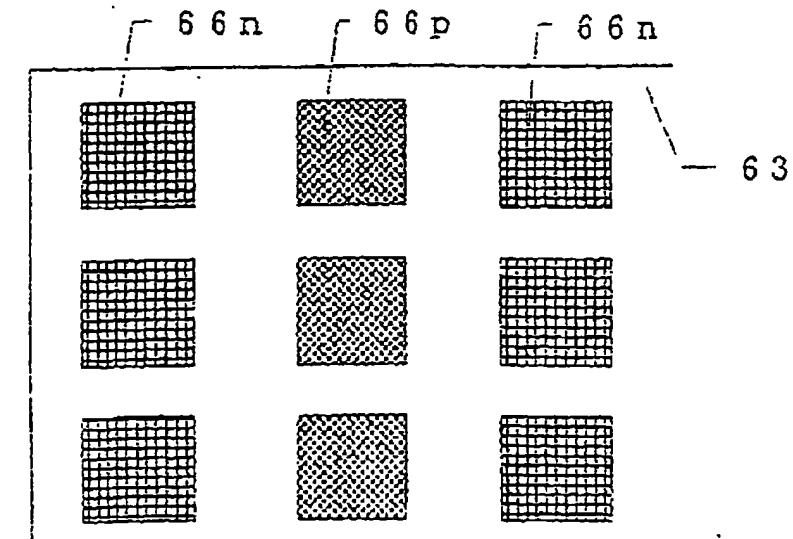
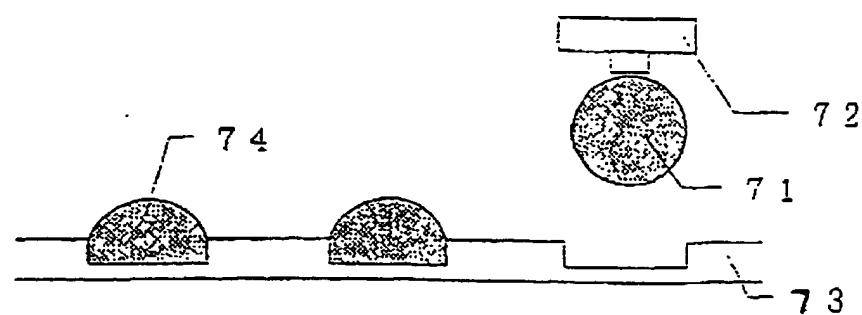
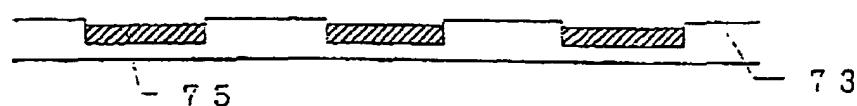


FIG. 7

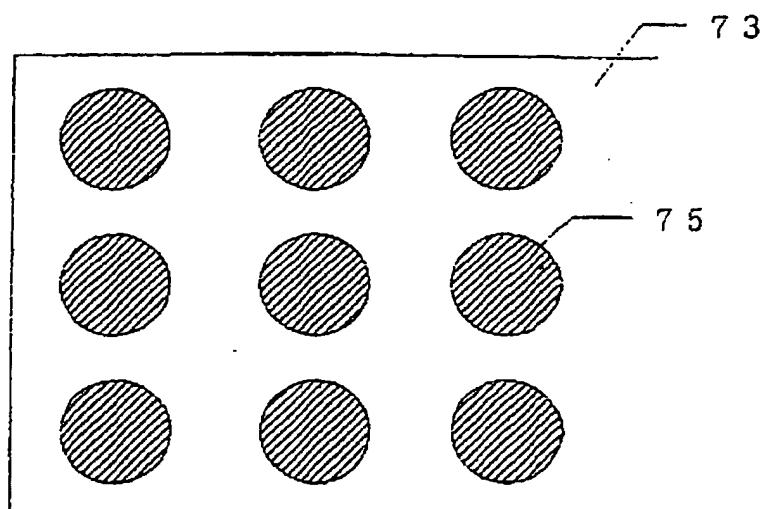
(a)



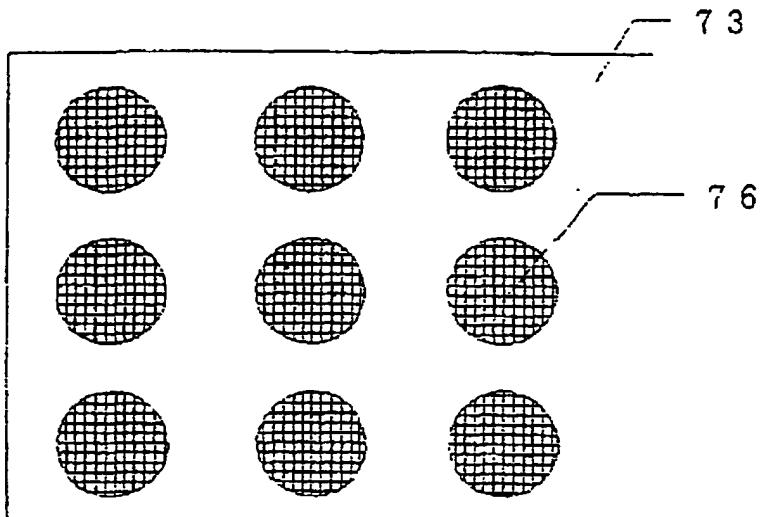
(b)



(c)



(d)



INTERNATIONAL SEARCH REPORT		International application No. PCT/JP00/01987																					
A. CLASSIFICATION OF SUBJECT MATTER Int.Cl ⁷ H01L21/208, C01B33/02																							
According to International Patent Classification (IPC) or to both national classification and IPC																							
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) Int.Cl ⁷ H01L21/208, 21/205, 21/20 Int.Cl ⁷ C01B33/02, 33/03, 33/04																							
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Jitsuyo shinan Koho 1926-1996 Jitsuyo Shinan Toroku Koho 1996-2000 Kokai Jitsuyo Shian Koho 1971-2000 Toroku Jitsuyo Shinan Koho 1994-2000																							
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)																							
C. DOCUMENTS CONSIDERED TO BE RELEVANT <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="text-align: left; padding-right: 20px;">Category*</th> <th style="padding-right: 20px;">Citation of document, with indication, where appropriate, of the relevant passages</th> <th style="padding-right: 20px;">Relevant to claim No.</th> </tr> </thead> <tbody> <tr> <td>X</td> <td>JP, 10-321536, A (Sharp Corporation), 04 December, 1998 (04.12.98), Full text (Family: none)</td> <td>14, 17, 19, 20</td> </tr> <tr> <td>Y</td> <td>JP, 3-215941, A (Hitachi Ltd.), 20 September, 1991 (20.09.91), Claims (Family: none)</td> <td>1-5, 8, 10, 11</td> </tr> <tr> <td>Y</td> <td>JP, 8-8179, A (Sharp Corporation), 12 January, 1996 (12.01.96), Full text (Family: none)</td> <td>4</td> </tr> <tr> <td>A</td> <td>JP, 11-79727, A (Sharp Corporation), 23 March, 1999 (23.03.99), Full text (Family: none)</td> <td>1-22</td> </tr> <tr> <td>A</td> <td>JP, 6-191821, A (Showa Denko K.K.), 12 July, 1994 (12.07.94), Full text (Family: none)</td> <td>1-22</td> </tr> <tr> <td>A</td> <td>JP, 5-144741, A (Showa Denko K.K.),</td> <td>1-22</td> </tr> </tbody> </table>			Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.	X	JP, 10-321536, A (Sharp Corporation), 04 December, 1998 (04.12.98), Full text (Family: none)	14, 17, 19, 20	Y	JP, 3-215941, A (Hitachi Ltd.), 20 September, 1991 (20.09.91), Claims (Family: none)	1-5, 8, 10, 11	Y	JP, 8-8179, A (Sharp Corporation), 12 January, 1996 (12.01.96), Full text (Family: none)	4	A	JP, 11-79727, A (Sharp Corporation), 23 March, 1999 (23.03.99), Full text (Family: none)	1-22	A	JP, 6-191821, A (Showa Denko K.K.), 12 July, 1994 (12.07.94), Full text (Family: none)	1-22	A	JP, 5-144741, A (Showa Denko K.K.),	1-22
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Date of the actual completion of the international search 20 June, 2000 (20.06.00)	Date of mailing of the international search report 04 July, 2000 (04.07.00)																						
Name and mailing address of the ISA/ Japanese Patent Office	Authorized officer																						
Facsimile No.	Telephone No.																						

INTERNATIONAL SEARCH REPORT		International application No. PCT/JP00/01987
C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
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	11 June, 1993 (11.06.93), Full text (Family: none)	

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